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Sheet 1 of 9

Complete if Known	
Application Number	10/042,203
Filing Date	January 11, 2002
First Named Inventor	Bernd RIEDL et al.
Group Art Unit	1625
Examiner Name	Rita J. Desai
Attorney Docket Number	BAYER-25A

U.S. PATENT DOCUMENTS

Examiner Initials*	Cite No.[1]	U.S. Patent Document Number	Kind Code[2] (if known)	Name of Patentee or Applicant of Cited Document	Date of Publication of Cited Document MM-DD-YYYY
RP	A1	2002/017507	A1	Santora et al.	11-21-2002
	A2	2002/0065283	A1	McMahon et al.	05-30-2002
	A3	2002/0065296	A1	Dumas et al.	05-30-2002
	A4	2004/0209905	A1	Kubo et al.	10-21-2004
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RP	A19	6608052	B2	Breitfelder et al.	08-19-2003

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RP	B1	WO	02/14311	A2	Amgen Inc.	02-21-2002		
	B2	WO	02/32872	A1	Eisai Co. Ltd.	04-25-2002		
	B3	WO	02/44158	A1	Pfizer Products Inc.	06-06-2002		
	B4	WO	02/07772	A2	Boehringer Ingelheim	01-31-2002		
	B5	WO	02/32872	A1	Eisai Co. Ltd.	04-25-2002		
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Examiner Signature	RJDesai	Date Considered	1/17/06

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	B26	WO	2005/048948	A2	Ambit Biosciences	06-02-2005		
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	B29	WO	02/085857	A2	Bayer Corporation	10-31-2002		
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Application Number	10/042,203
Filing Date	January 11, 2002
First Named Inventor	Bernd RIEDL et al.
Group Art Unit	1625
Examiner Name	Rita J. Desai
Attorney Docket Number	BAYER-25A

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RD	B38	WO	01/07411	A1	Banyu Pharmaceutical	02-01-2001		
RD	B39	UK	1110099		Julia	06-07-1966		
RD	B40	WO	2004078746	A2	Boyer et al.	09-16-2004		
RD	B41	WO	2005037273	A1	Ramurthy et al.	04-28-2005		
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Sheet 4 of 9

Complete if Known	
Application Number	10/042,203
Filing Date	January 11, 2002
First Named Inventor	Bernd RIEDL et al.
Group Art Unit	1625
Examiner Name	Rita J. Desai
Attorney Docket Number	BAYER-25A

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NON PATENT LITERATURE DOCUMENTS

Examiner Initials*	Cite No.[1]	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T[2]
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Examiner Signature	R Desai	Date Considered	1/17/06

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Sheet 5 of 9

Complete if Known	
Application Number	10/042,203
Filing Date	January 11, 2002
First Named Inventor	Bernd RIEDL et al.
Group Art Unit	1625
Examiner Name	Rita J. Desai
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Examiner Signature	R Desai	Date Considered	1/17/06

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Group Art Unit	1625
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	C29	Heim et al., "Antitumor effect and potentiation or reduction in cytotoxic drug activity in human colon carcinoma cells by the Raf kinase inhibitor (RKI) BAY 43-9006," *International Journal of Clinical Pharmacology and Therapeutics*, Vol. 41, No. 12/2003 (616-617)	
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	C31	Mross et al., "Drug-drug reaction pharmacokinetic study with the Raf kinase inhibitor (RKI) BAY 43-9006 administered in combination with irinotecan (CPT-11) in patients with solid tumors," *International Journal of Clinical Pharmacology and Therapeutics*, Vol. 41, No. 12/2003 (618-619)	
RD	C32	Richly et al., "A phase I clinical and pharmacokinetic study of the Raf kinase inhibitor (RKI) BAY 43-9006 administered in combination with doxorubicin in patients with solid tumors," *International Journal of Clinical Pharmacology and Therapeutics*, Vol. 41, No. 12/2003 (620-621)	

Examiner Signature	RDesai	Date Considered	1/17/06

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Filing Date	January 11, 2002
First Named Inventor	Bernd RIEDL et al.
Group Art Unit	1625
Examiner Name	Rita J. Desai
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RP	C33	DeGrendele, "Activity of the raf kinase inhibitor BAY 43-9006 in patients with advanced solid tumors," *Clinical Colorectal Cancer*, May 2003, pp. 16-18	
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	C35	Thompson et al., "Recent progress in targeting the Raf/MEK/ERK pathway with inhibitors in cancer drug discovery," *Curr. Opin. Pharmacol.*, 2005 Aug., 5(4):350-6	
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	C41	Regan et al., "Pyrazole urea-based inhibitors of p38 MAP kinase: from lead compound to clinical candidate," *J. Med. Chem.*, 2002, 45, 2994-3008	
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	C45	Jeffcoat et al., "The metabolism and toxicity of halogenated carbanilides," *Drug Metabolism and Deposition*, Vol. 5, No. 2, 157-166	
RP	C46	Murata et al., "Facile synthesis of new pyrrolo[3,4-*d*]pyrimidine-2,4-diones," *Chem. Pharm. Bull.*, 22(5) 1212-1213 (1974)	

Examiner Signature	RJDesai	Date Considered	1/17/06

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Sheet 8 of 9

Complete If Known	
Application Number	10/042,203
Filing Date	January 11, 2002
First Named Inventor	Bernd RIEDL et al.
Group Art Unit	1625
Examiner Name	Rita J. Desai
Attorney Docket Number	BAYER-25A

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RP	C47	Yasuo et al, "Intra-arterial ACNU, CDDP chemotherapy for brain metastases from lung cancer: comparison of cases with and without intra-arterial mannitol infusion," *Neurol. Surg.*, (1993) vol. 21, no. 6, pp. 513-518	
	C48	Murata et al., "Facile synthesis of new pyrrolo[3,4-d]pyrimidine-2,4-diones," *Chemical & Pharmaceutical Bulletin*, (1974), 22(5):1212-13	
	C49	Hanson, "Inhibitors of p38 kinase," *Expert Opinion on Therapeutic Patents*, July 1997, vol. 7, no. 7, pp. 729-733(5)	
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	C53	Dumas, "Protein kinase inhibitors from the ureas class," *Current Opinion in Drug Discovery & Development*, 2002, Vol. 5, No. 5, 718-727	
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	C56	Wisaner et al., "Analogues of platelet activating factor. 7. Bis-aryl amide and bis-aryl urea receptor antagonists of PAF," *J. Med. Chem.*, 1992, 35, 4779-4789	
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	C58	Lemoine, "Overview of ras oncogenes and their clinical potential," Chapter 10,	
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	C60	Siu et all, "Phase I study of oral raf-1 kinase inhibitor BAY 43-9006 with gemcitabine in patients with advanced solid tumors," *Proc Am Soc Clinic Oncol*, 22:207, 2003 (abstr 828)	
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RD	C63	Adjei et al., "A phase I study of BAY 43-9006 and gefitinib in patients with refractory or recurrent non-small-cell lung cancer (NSCLC)," Meeting: *2005 ASCO Annual Meeting*, Category: Developmental Therapeutics: Molecular Therapeutics, Subcategory: Antianglogenic or Antimetastatic agents, Abstract No. 4510	
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	C65	Van Muijlwijk-Koezen et al., "Isoquinoline and quinazoline urea analogues as antagonists for the human adenosine A_3 receptor," *J. Med. Chem.*, 2000, 43, 2227-2238	
	C66	Eisenhauer et al., "Impact of new non-cytotoxics in the treatment in ovarian cancer," *Int. J. Gynecol Cancer*, 2001, 11 (Suppl. 1), 68-72	
	C67	Kubo et al., "Synthesis and structure-activity relationship of quinazoline-urea derivatives as novel orally active VEGF receptor tyrosine kinase selective inhibitors," #913, XP-001152608	
	C68	Carter et al, "Anti-tumor efficacy of the orally active raf kinase inhibitor BAY 43-9006 in human tumor xenograft models," #4954, XP-001145482	
	C69	Strumberg et al., "Phase I and pharmacokinetic study of the raf kinase inhibitor bay 43-9006 in patients with locally advanced or metastatic cancer," #2921, XP-001145481	
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	C72	Iwadate et al., "Intra-arterial ACNU, CDDP chemotherapy for brain metastases from lung cancer: comparison of cases with and without intra-arterial mannitol infusion," Dept of Neurological Surgery, Chiba Cancer Center Hospital, Clinical Trial, Journal Article, Randomized Controlled Trial, Vol. 21, No. 6, 513-518	
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RD	C74	Cunningham et al., "A phase I trial of H-ras antisense oligonucleotide ISIS 2503 administered as a continuous intravenous infusion in patients with advanced carcinoma," *Cancer*, September 2001, Vol. 92, No. 5, 1265-1271	

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